DOCKET NO.: 48378-0003-00-&S

Application No.: 10/621,711

Office Action Dated: January 18, 2007

PATENT REPLY FILED UNDER EXPEDITED PROCEDURE PURSUANT TO 37 CFR § 1.116

This listing of claims will replace all prior versions, and listings, of claims in the application.

Listing of Claims:

1-162. (Canceled)

- 163. (Previously presented) A transdermal delivery system comprising a backing layer and an adhesive polymer matrix affixed to the backing layer, wherein the adhesive polymer matrix comprises:
 - (a) an adhesive polymer;
 - (b) a humectant;
 - (c) a combination of skin permeation enhancing agents comprising dimethyl sulfoxide; a fatty (C₈-C₂₀) alcohol ester of lactic acid; a lower (C₁-C₄) alkyl ester of lactic acid; and capric acid;
 - (d) a progestin; and
 - (e) an estrogen.
- 164. (Previously presented) The transdermal delivery system of claim 163, wherein the adhesive polymer is a polyacrylate copolymer, a polyisobutylene or a silicone adhesive.
- 165. (Previously presented) The transdermal delivery system of claim 164, wherein the polyacrylate copolymer comprises a 2-ethylhexyl acrylate monomer.
- 166. (Previously presented) The transdermal delivery system of claim 165, wherein the polyacrylate copolymer further comprises about 3 to 60% w/w vinyl acetate.
- 167. (Previously presented) The transdermal delivery system of claim 163, wherein the humectant comprises polyvinylpyrrolidone.
- 168. (Previously presented) The transdermal delivery system of claim 167, wherein the humectant comprises a polyvinylpyrrolidone copolymer.

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169. (Previously presented) The transdermal delivery system of claim 168, wherein the humectant is a polyvinylpyrrolidone/vinyl acetate copolymer.

- 170. (Previously presented) The transdermal delivery system of claim 169, wherein the polyvinylpyrrolidone is formulated in an amount of about 60% w/w and the vinyl acetate is formulated in an amount of about 40% w/w in the polyvinylpyrrolidone/vinyl acetate copolymer.
- 171. (Previously presented) The transdermal delivery system of claim 163, wherein the fatty alcohol ester of lactic acid is lauryl lactate.
- 172. (Previously presented) The transdermal delivery system of claim 163, wherein the lower alkyl ester of lactic acid is ethyl lactate.
- 173. (Previously presented) The transdermal delivery system of claim 163, wherein the progestin is levonorgestrel.
- 174. (Previously presented) The transdermal delivery system of claim 163, wherein the estrogen is ethinyl estradiol or 17 β -estradiol.
- 175. (New) The transdermal delivery system of claim 173, which, when applied to the skin of a human, once each week, consecutively over a period of three or more weeks, deliver *in vivo* an average serum concentration of over 1000 pg/ml of levonorgestrel.